


**UNITED STATES ENVIRONMENTAL PROTECTION AGENCY**
**401 M Street, S.W.  
WASHINGTON, D.C. 20460**

**Paperwork Reduction Act Notice:** The public reporting burden for this collection of information is estimated to average 1.25 hours per response for registration activities and 0.25 hours per response for reregistration and special review activities, including time for reading the instructions and completing the necessary forms. Send comments regarding the burden estimate or any other aspect of this collection of information, including suggestions for reducing the burden to: Director, OPPE Information Management Division (2137), U.S. Environmental Protection Agency, 401 M Street, S.W., Washington, DC 20460. Do not send the form to this address.

**Certification with Respect to Citation of Data**

<b>Applicant's/Registrant's Name, Address and Telephone Number</b> Nufarm Limited C/O Nufarm Americas Inc., Agent for Nufarm Limited 4020 Aerial Center Parkway, Suite 101, Morrisville, NC 27560	<b>EPA Registration Number/ File Symbol</b>  35935-XXX
<b>Active Ingredient(s) and/or representative test compound(s):</b> Mefenoxam	<b>Date</b> March 22, 2019
<b>General use pattern(s) (list all those claimed for this product using 40 CFR Part 158)</b> Terrestrial food crops; Terrestrial feed crops; Terrestrial nonfood crops; Greenhouse nonfood crops; Residential outdoor use	<b>Product Name</b> Nufarm Mefenoxam Technical

**NOTE:** If your product is a 100% repackaging of another purchased EPA-registered product labeled for all the same uses on your label, you do not need to submit this form. You must submit the Formulator's Exemption Statement (EPA Form 8570-27).

☐ I am responding to a Data Call-In Notice, and have included with this form a list of companies sent offers of compensation (the Data Matrix form should be used for this purpose).

**SECTION I: METHOD OF DATA SUPPORT (Check one method only)**

☐ I am using the cite-all method of support, and have included with this form a list of companies sent offers of compensation (the Data Matrix form should be used for this purpose).

☒ I am using the selective method of support (or cite-all option under the selective method), and have included with this form a completed list of data requirements (the Data Matrix form must be used).

**SECTION II: GENERAL OFFER TO PAY**

[Required if using the cite-all method or when using the cite-all option under the selective method to satisfy one or more data requirements]

☒ I hereby offer and agree to pay compensation, to other persons, with regard to the approval of this application, to the extent required by FIFRA.

**SECTION III: CERTIFICATION**

I certify that this application for registration, this form for reregistration, or this Data Call-In Notice is supported by all data submitted or cited in the application for registration, the form for reregistration, or this Data Call-In response. In addition, if cite-all option under the selective method is indicated in Section I, this application is supported by all data in the Agency's files that (1) concern the properties or effects of this product or an identical or substantially similar product, or one or more of the ingredients in this product; and (2) is a type of data that would be required to be submitted under the data requirements in effect on the date of approval of this application if the application sought the initial registration of a product of identical or similar composition and uses.

I certify that for each exclusive use study cited in support of this registration or reregistration, that I am the original submitter or that I have obtained the written permission of the original submitter to cite that study.

I certify that for each study cited in support of this registration or reregistration that is not an exclusive use study, either: (a) I am the original data submitter; (b) I have obtained the written permission of the original data submitter to use this study in support of this application; (c) all periods of eligibility for compensation have expired for the study; (d) the study is in the public literature; or (e) I have notified in writing the company that submitted the study and have offered (i) to pay compensation to the extent required by sections 3(c)(1)(F) and/or 3(c)(2)(B) of FIFRA; and (ii) to commence negotiations to determine the amount and terms of compensation, if any, to be paid for the use of the study.

I certify that in all instances where an offer of compensation is required, copies of all offers to pay compensation and evidence of their delivery in accordance with sections 3(c)(1)(F) and/or 3(c)(2)(B) of FIFRA are available and will be submitted to the agency upon request. Should I fail to produce such evidence to the Agency upon request, I understand that the Agency may initiate action to deny, cancel or suspend the registration of my product in conformity with FIFRA.

**I certify that the statements I have made on this form and all attachments to it are true, accurate and complete. I acknowledge that any knowingly false or misleading statements may be punishable by fine or imprisonment or both under applicable law.**

<b>Signature</b> 	<b>Date</b> March 22, 2019	<b>Typed or Printed Name and Title</b> Danielle A. Larochelle Sr. Regulatory Manager
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<b>Active Ingredient(s) and/or representative test compound(s):</b> Mefenoxam	<b>Date</b> September 16, 2021
<b>General use pattern(s) (list all those claimed for this product using 40 CFR Part 158)</b> Terrestrial food crops; Terrestrial feed crops; Terrestrial nonfood crops; Greenhouse nonfood crops; Residential outdoor use	<b>Product Name</b> Nufarm Mefenoxam Technical

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